Impaired Increase of Retinal Capillary Blood Flow to Flicker Light Exposure in Arterial Hypertension

Martin Ritt, Joanna M. Harazny, Christian Ott, Ulrike Raff, Philipp Bauernschubert, Marina Lehmann, Georg Michelson, Roland E. Schmieder

Abstract—We hypothesized that the increase of retinal capillary blood flow (RCF) to flicker light exposure is impaired in subjects with arterial hypertension. In 146 nondiabetic untreated male subjects with (n=50) or without (n=96) arterial hypertension, RCF was measured before and after flicker light exposure noninvasively and in vivo using scanning laser Doppler flowmetry. In addition, in a subgroup of 28 subjects, the change of RCF to flicker light exposure was again assessed during parallel infusion of nitric oxide synthase inhibitor N-monomethyl-L-arginine (L-NMMA). The increase of RCF to flicker light exposure was lower in patients with untreated hypertension compared with normotensive subjects when expressed in absolute terms (7.69±5.4 versus 27.2±4.4 AU; \( P = 0.013 \)) or percent changes (2.95±4 versus 8.33±12%; \( P = 0.023 \)). Systolic (\( \beta = -0.216; P = 0.023 \)) but not diastolic blood pressure (\( \beta = -0.117; P = 0.243 \)) or mean arterial pressure (\( \beta = -0.178; P = 0.073 \)) was negatively related to the percent change of RCF to flicker light exposure, independently of other cardiovascular risk factors. In the subgroup of 28 subjects, the increase of RCF to flicker light exposure was similar at baseline and during parallel infusion of L-NMMA when expressed in absolute terms (20.0±51 versus 22.6±56 AU; \( P = 0.731 \)) or percent changes (7.12±16 versus 8.29±18%; \( P = 0.607 \)). The increase of RCF to flicker light exposure is impaired in arterial hypertension. In the subgroup of the total study cohort, nitric oxide was not a major determinant of the increase of RCF to flicker light exposure. (Hypertension. 2012;60:871-876.)

Key Words: retina ▪ capillary blood flow ▪ vasodilatory properties ▪ nitric oxide ▪ arterial hypertension

Arterial hypertension is a major determinant of morbidity and mortality due to cardiovascular complications. \(^1\) Reduction in systolic and diastolic blood pressure levels was found to reduce and slow the occurrence of cardiovascular events in subjects with arterial hypertension; \(^2\) however, blood pressure levels per se might be unreliable indicators of cardiovascular risk in the individual subjects. In arterial hypertension, elevated blood pressure levels lead to structural and functional changes of blood vessels and organ damage, such as retinopathy, thickening of carotid arteries, large artery stiffening, left ventricular hypertrophy, and increased urinary albumin excretion, among others. These subclinical parameters represent intermediate end points that frequently precede major cardiovascular events and indicate the need for aggressive medical blood pressure control in context of individual global cardiovascular risk profile. \(^2\)

Since the famous work by Keith, Wagener, and Barker, \(^3\) several studies have demonstrated the prognostic significance of retinal vascular alterations for predicting morbidity and mortality in subjects with arterial hypertension; \(^4,6\) however, owing to improvement in patient management, nowadays, grade 3 and grade 4 hypertensive retinopathy are seldom observed, and grade 1 and 2 hypertensive retinopathy reveal low power for predicting cardiovascular events, the classical classification system of hypertensive retinopathy, dating back to the aforementioned work, \(^9\) has repeatedly been criticized in the last 2 decades. \(^7,8\) In parallel, much research effort has focused on the evaluation of early retinal alterations in arterial hypertension as potential new parameters of hypertensive target organ damage and retinopathy. \(^9-11\)

Impairment of peripheral vasodilatory properties \(^12,13\) represent early vascular changes in arterial hypertension. Of clinical interest, impaired peripheral vasodilatory properties was found to reveal prognostic significance, with respect to adverse cardiovascular outcome in subjects with arterial hypertension. \(^14\) Whether vasodilatory properties are impaired in the retinal vascular bed in subjects with arterial hypertension has not yet been examined. Improvement in imaging technology nowadays allows assessment of retinal capillary blood flow noninvasively and in vivo in humans. Moreover, exposure to flicker light was found to increase capillary blood flow in the retinal circulation. \(^15\) The mechanisms for the increase of retinal capillary blood flow to flicker light are incompletely understood, but data in healthy subjects indicate that nitric oxide might (at least, in part) be involved. \(^16\)

In the current study, we hypothesized that the increase of retinal capillary blood flow to flicker light is impaired in

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subjects with arterial hypertension. Moreover, we aimed to assess the contribution of nitric oxide on the flicker light-induced increase of retinal capillary blood flow in a subgroup of the study cohort.

Methods

Study Design and Study Population

This observational study was performed at the Clinical Research Unit of the Department of Nephrology and Hypertension, University of Erlangen-Nuremberg, Germany. Study participants were recruited through advertisements in local newspapers. Patients with arterial hypertension (defined as systolic blood pressure \(\geq 140\) mm Hg and/or diastolic blood pressure \(\geq 90\) mm Hg) and normotensive individuals (systolic blood pressure \(<140\) mm Hg and diastolic blood pressure \(<90\) mm Hg) of male gender, between 18 and 75 years of age, were included. Exclusion criteria were history or any present clinical evidence for cardiovascular disease; atrial fibrillation or atrioventricular blockade, grade II or higher; history or current use of any antihypertensive drug or other medication; renal impairment (defined as estimated creatinine-clearance \(<60\) mL/min according to the formula by Cockroft and Gault); hepatic disease, diabetes mellitus (defined by fasting glucose \(\geq 126\) mg/dL or on glucose-lowering medication or history of diabetes mellitus); any form of secondary arterial hypertension; any significant eye disease (including hypertensive retinopathy, grade III and IV), and smoking. Blood pressure was measured in a sitting position, according to World Health Organization criteria, 3%, and the average was calculated. Before enrolment in the study, written informed consent was obtained from each participant. The study protocol was approved by the Clinical Investigations Ethics Committee of the University Erlangen-Nuremberg. The study was conducted in accordance with Good Clinical Practice guidelines and in compliance with the Declaration of Helsinki.

Assessment of Parameters of Retinal Vessels

Measurement of Retinal Capillary Blood Flow

Retinal capillary blood flow was assessed using scanning laser Doppler flowmetry at 670 nm (Heidelberg Engineering). Briefly, a retinal sample of length 2.56 mm (256 points)\(^/\text{mm}\) was measured.\(^{17}\) Measurements were performed in the juxtapapillary area of the right eye, 2 to 3 mm temporally to the optic nerve; the average from 3 singular measurements was taken. The confocal technique of the device returned to baseline levels (data not shown).

Baseline Clinical Characteristics

Baseline Clinical Characteristics

The characteristics of the study participants, stratified into subjects with arterial hypertension and normotensive controls, are given in Table 1: Subjects with arterial hypertension were older and revealed higher body weight, body mass index, blood pressure levels, heart rate, fasting glucose, and triglyceride levels compared with normotensive patients. Height, high-density lipoprotein (HDL)-cholesterol, low-density lipoprotein (LDL)-cholesterol, serum creatinine, and
estimated creatinine clearance did not differ between the 2 groups.

Baseline Retinal Characteristics and Changes of Retinal Capillary Blood Flow to Flicker Light Exposure
The baseline retinal characteristics and changes in retinal parameter to flicker light exposure of the study participants, stratified into patients with arterial hypertension and normotensive controls, are also given in Table 1: Retinal capillary blood flow at baseline (ie, before flicker light exposure) did not differ between the 2 patient groups. The increase of retinal capillary blood flow to flicker light exposure was lower in subjects with arterial hypertension than in normotensive controls, irrespective whether expressed in absolute terms or in percent changes, even after adjustment of the analysis for possible confounders.

Univariate Correlation Analysis Between Various Clinical Parameters and the Percent Change of Retinal Capillary Blood Flow to Flicker Light Exposure
Systolic blood pressure (Figure A) and mean arterial pressure ($r = -0.171; P = 0.039$) were negatively related to the percent increase of retinal capillary blood flow to flicker light exposure. The negative relationship of diastolic blood pressure to the percent increase of retinal capillary blood flow to flicker light exposure did not reach statistical significance (Figure B). None of the other clinical parameters, including age ($r = 0.002; P = 0.980$), height ($r = 0.115; P = 0.168$), weight ($r = 0.080; P = 0.337$), body mass index ($r = 0.047; P = 0.572$), heart rate ($r = -0.008; P = 0.924$), fasting glucose ($r = -0.005; P = 0.956$), triglyceride levels ($r = -0.053; P = 0.529$), LDL-cholesterol ($r = -0.114; P = 0.175$), HDL-cholesterol ($r = -0.007; P = 0.229$), and serum creatinine clearance ($r = 0.007; P = 0.933$) and retinal capillary blood flow at baseline (ie, before flicker light exposure) ($r = -0.150; P = 0.072$) revealed a significant relationship to the percent increase of retinal capillary blood flow to flicker light.

Multiple Linear Regression Analysis Assessing the Quantitative Role of Major Cardiovascular Risk Factors in Relation to Each Other on the Percent Change of Retinal Capillary Blood Flow to Flicker Light Exposure
Systolic (model 1) but not diastolic blood pressure (model 2) or mean arterial pressure (model 3) was negatively related to the percent increase of retinal capillary blood flow to flicker light exposure, independently of other cardiovascular risk factors (see Table 2). None of the other cardiovascular risk factors revealed such an independent relationship to the percent increase of retinal capillary blood flow to flicker light exposure (see Table 2).

### Table 1. Clinical and Retinal Characteristics of the Study Cohort, Stratified Into Subjects With Arterial Hypertension and Normotensive Controls

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Hypertensive Subjects (n=50)</th>
<th>Normotensive Subjects (n=96)</th>
<th>P Value</th>
<th>P Value Adjusted*†</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Clinical characteristics</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (y)</td>
<td>40.5±7.8</td>
<td>36.8±8.9</td>
<td>0.011</td>
<td>...</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>182±6.5</td>
<td>182±6.0</td>
<td>0.867</td>
<td>...</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>99.5±16</td>
<td>92.2±18</td>
<td>0.014</td>
<td>...</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>30.0±4.9</td>
<td>27.8±5.0</td>
<td>0.012</td>
<td>...</td>
</tr>
<tr>
<td>Systolic blood pressure (mm Hg)</td>
<td>146±8.6</td>
<td>129±6.4</td>
<td>&lt;0.001</td>
<td>...</td>
</tr>
<tr>
<td>Diastolic blood pressure (mm Hg)</td>
<td>88.4±7.0</td>
<td>76.5±6.4</td>
<td>&lt;0.001</td>
<td>...</td>
</tr>
<tr>
<td>Mean arterial pressure (mm Hg)</td>
<td>108±6.2</td>
<td>93.8±5.2</td>
<td>&lt;0.001</td>
<td>...</td>
</tr>
<tr>
<td>Heart rate (beats/minute)</td>
<td>75.5±9.9</td>
<td>71.2±10</td>
<td>0.015</td>
<td>...</td>
</tr>
<tr>
<td>Fasting glucose (mg/dl)</td>
<td>101±16</td>
<td>95.5±13</td>
<td>0.028</td>
<td>...</td>
</tr>
<tr>
<td>Triglycerides (g/dl)</td>
<td>199±121</td>
<td>159±90</td>
<td>0.027</td>
<td>...</td>
</tr>
<tr>
<td>HDL-cholesterol (mg/dl)</td>
<td>50.4±13</td>
<td>49.2±8.7</td>
<td>0.564</td>
<td>...</td>
</tr>
<tr>
<td>LDL-cholesterol (mg/dl)</td>
<td>152±34</td>
<td>143±39</td>
<td>0.163</td>
<td>...</td>
</tr>
<tr>
<td>Serum creatinine (mg/dl)</td>
<td>0.9±0.1</td>
<td>0.89±0.1</td>
<td>0.212</td>
<td>...</td>
</tr>
<tr>
<td>Estimated creatinine clearance (ml/min/1.73 m²)</td>
<td>118±22</td>
<td>120±20</td>
<td>0.496</td>
<td>...</td>
</tr>
<tr>
<td><strong>Retinal characteristics</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Retinal capillary blood flow at baseline (ie, before flicker light exposure) (AU)</td>
<td>360±89</td>
<td>334±83</td>
<td>0.077</td>
<td>0.229</td>
</tr>
<tr>
<td>Change of retinal capillary blood flow to flicker light exposure (AU)</td>
<td>7.69±54</td>
<td>27.2±44</td>
<td>0.021</td>
<td>0.012/0.013</td>
</tr>
<tr>
<td>Change of retinal capillary blood flow to flicker light exposure (%)</td>
<td>2.95±14</td>
<td>8.33±12</td>
<td>0.019</td>
<td>0.016/0.023</td>
</tr>
</tbody>
</table>

*P adjusted for variables that differed with a P value of <0.05 between the hypertensive and normotensive group (ie, age, weight, body mass index, heart rate, fasting glucose, and triglyceride levels).
†P adjusted for variables that differed with a P value of <0.10 between the hypertensive and normotensive group (ie, age, weight, body mass index, heart rate, fasting glucose, triglyceride levels, and retinal capillary blood flow at baseline (ie, before flicker light exposure).
Assessment of the Contribution of Nitric Oxide on the Increase of Retinal Capillary Blood Flow to Flicker Light Exposure in a Subgroup (n=28) of the Total Study Cohort

The clinical and retinal characteristics of these 28 subjects were as follows: age, 38.9±5.5 years; height, 181±6.3 cm; weight, 110±12 kg; body mass index, 33±3.9 kg/m²; systolic blood pressure, 135±10 mm Hg; diastolic blood pressure, 81±7.9 mm Hg; mean arterial pressure, 98.8±7.8 mm Hg; heart rate, 74.8±9.1 beats per minute; fasting glucose, 101±23 mg/dL; triglycerides, 171±77 mg/dL; HDL-cholesterol, 48.6±11 mg/dL; LDL-cholesterol, 150±36 mg/dL; serum creatinine, 0.92±0.1 mg/dL; estimated creatinine clearance, 126±21 mL/min/1.73 m²; and retinal capillary blood flow, 337±56 AU.

The increase of retinal capillary blood flow to flicker light exposure did not differ compared with the increase in retinal capillary blood flow to flicker light exposure during infusion of L-NMMA when expressed in absolute terms (18%; 16 versus 8.29 ± 7.877 g/dL; HDL-cholesterol, 23 mg/dL; triglycerides, 171 mg/dL; LDL-cholesterol, 150±36 mg/dL; serum creatinine, 0.92±0.1 mg/dL; estimated creatinine clearance, 126±21 mL/min/1.73 m²; and retinal capillary blood flow, 337±56 AU.

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The underlying mechanisms explaining the impaired increase of retinal capillary blood flow to flicker light exposure in our patients with hypertension compared with normotensive controls remains unclear. Reduced production and/or action of vasorelaxing factors such as nitric oxide, endothelial-derived hyperpolarizing factor, and prostacyclin or increased production or activity of vasoconstrictor factors such as prostanoids (endoperoxides, thromboxane A2), superoxide anions, and endothelin increases vascular tone, leads to chronic vasoconstriction of peripheral vessels, and might impair vasodilatory properties in arterial hypertension.

The major finding of our study is that the increase of retinal capillary blood flow to flicker light exposure (%)

Discussion

The major finding of our study is that the increase of retinal capillary blood flow to flicker light exposure is impaired in subjects with arterial hypertension compared with normotensive controls. Moreover, systolic but not diastolic blood pressure or mean arterial pressure was found to be negatively related to the percent increase of retinal capillary blood flow to flicker light exposure, independently of other cardiovascular risk factors. Thus, the assessment of changes of retinal capillary blood flow to flicker light exposure might be an interesting noninvasive in vivo tool for detection of early vascular changes in arterial hypertension.

It might be rational to hypothesize that either constricted vessels with increased vascular resistance in arterial hypertension might be associated with decreased capillary blood flow and to suggest that changes in baseline capillary blood flow before the flicker light exposure may result in altered relative changes in capillary blood flow after the flicker light exposure; however, in our study, retinal capillary blood flow at baseline (ie, before flicker light exposure) did not differ significantly between the hypertensive and the normotensive group. Moreover, neither in univariate correlation analysis nor in multiple linear regression analysis, a significant relationship between retinal capillary blood flow at baseline and the increase of retinal capillary blood flow to flicker light exposure was found.

The underlying mechanisms explaining the impaired increase of retinal capillary blood flow to flicker light exposure in our patients with hypertension compared with normotensive controls remains unclear. Reduced production and/or action of vasorelaxing factors such as nitric oxide, endothelial-derived hyperpolarizing factor, and prostacyclin or increased production or activity of vasoconstrictor factors such as prostanoids (endoperoxides, thromboxane A2), superoxide anions, and endothelin increases vascular tone, leads to chronic vasoconstriction of peripheral vessels, and might impair vasodilatory properties in arterial hypertension.

Figure A, Relationship between the percent change of retinal capillary blood flow to flicker light exposure and systolic blood pressure. B, Relationship between the percent change of retinal capillary blood flow to flicker light exposure and diastolic blood pressure.
Characteristics under model 1, 2, and 3 are characteristics that were included in that particular model.

increase of retinal capillary blood flow to flicker light. This result does not support the hypothesis of a major impact of nitric oxide on the increase of retinal capillary blood flow to flicker light exposure at first glance; however, in contrast to our study, Dorner et al\textsuperscript{16} reported a reduced increase of major retinal arteriolar diameters in response to flicker light exposure during concomitant infusion of 3 mg/kg body weight of L-NMMA, a dosage comparable to the dosage used in our current study, in a cohort of 12 young (mean ± SD age of 25.6 ± 2.1 years) healthy, nonsmoking subjects. The reasons for the discrepancy between the 2 studies might be owing to differences in the methodologies used: Dorner et al\textsuperscript{16} measured changes in major retinal arteriolar diameter in reflection images (by digital retinal photography), while we measured true changes in perfusion of the retinal capillary bed by scanning laser Doppler flowmetry. Second, Dorner et al\textsuperscript{16} used a frequency of the flicker light of 8 Hz, while we used a frequency of flicker light of 10 Hz. Previous data indicate that the frequency of flicker light might indeed impact on retinal capillary blood flow response to flicker light exposure.\textsuperscript{27} Third, Dorner et al\textsuperscript{16} used a much shorter flicker light duration of up to 64 seconds compared with a 3-minute flicker period in our current study. Furthermore, most of the patients in our substudy revealed cardiovascular risk factors such as arterial hypertension, metabolic syndrome, or obesity that might have resulted in decreased basal nitric oxide activity.\textsuperscript{20} Thus, to clearly address the role of nitric oxide in the increase of retinal capillary blood flow to flicker light exposure under physiological conditions, a cohort of healthy volunteers who do not reveal any cardiovascular risk factors will need to be examined.

Our study has limitations. First, although the readers of retinal vasculature were blinded to the clinical data (including blood pressure levels and history of arterial hypertension, among others), they might have been biased by detecting signs of hypertensive retinopathy, such as focal arteriolar narrowing among others; however, such signs of hypertensive retinopathy might also be detectable in patients with prehypertension\textsuperscript{7} that, in case of the current study, have been grouped into the normotensive group. Second, owing to inclusion criteria, our study cohort comprised only male subjects. Hormone levels, that potentially impact on microvascular function, differ between women and men.\textsuperscript{28} Therefore, results of our study might not be extensible to women. Third, all subjects in our current study were white. It cannot be excluded that the impact of blood pressure on the flicker light-induced changes in retinal capillary blood flow might differ between different ethnic groups. Fourth, owing to the cross-sectional design of our study, we can only describe associations but cannot give any information with respect to a causal role.

Perspectives
We found that the increase of retinal capillary blood flow to flicker light is impaired in subjects with arterial hypertension. Systolic but not diastolic blood pressure or mean arterial pressure was negatively related to the increase of retinal capillary blood flow to flicker light, independently of other cardiovascular risk factors. Thus, the noninvasive measurement of increases of retinal capillary blood flow to flicker light exposure might represent an interesting research and potentially clinical tool to assess early microvascular changes in vivo in arterial hypertension.

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Disclosures
None.
References


Novelty and Significance

What Is New?

- The increase of retinal capillary blood flow to flicker light exposure, assessed by scanning laser Doppler flowmetry, is lower in hypertensive compared to normotensive subjects and negatively related to systolic blood pressure, independently of possible confounders.

What Is Relevant?

- Assessment of changes in retinal capillary blood flow to flicker light exposure by scanning laser Doppler flowmetry allows detection of early microvascular changes in hypertension.
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